

Structured Training and Staff Retention: A 3-year review

Ashley Anderson, RN, MSN, CRNP; Hilary Avery, RN, BSN; Diana Hablitz, RN, BSN, CRNP, CCRP; Weiping DeBlasio, RN, BSN, MBA
O'Neal Comprehensive Cancer Center at the University of Alabama at Birmingham

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Background

High turnover among clinical research staff—including Clinical Research Coordinators, Data Coordinators, and Regulatory Coordinators—poses significant risks to study quality, protocol compliance, and operational continuity. This presentation examines how structured training and education programs can enhance workforce stability within clinical research settings. Our study was conducted retrospectively over the course of a 3-year review.

Goals

A structured training program was implemented due to an increasing rate of staff turnover. In early 2023, the Education Department in the Cancer Center set the goal of creating a structured training program to assist in the stabilization of staff turnover and provide career development pathways for Clinical Trials Office staff.

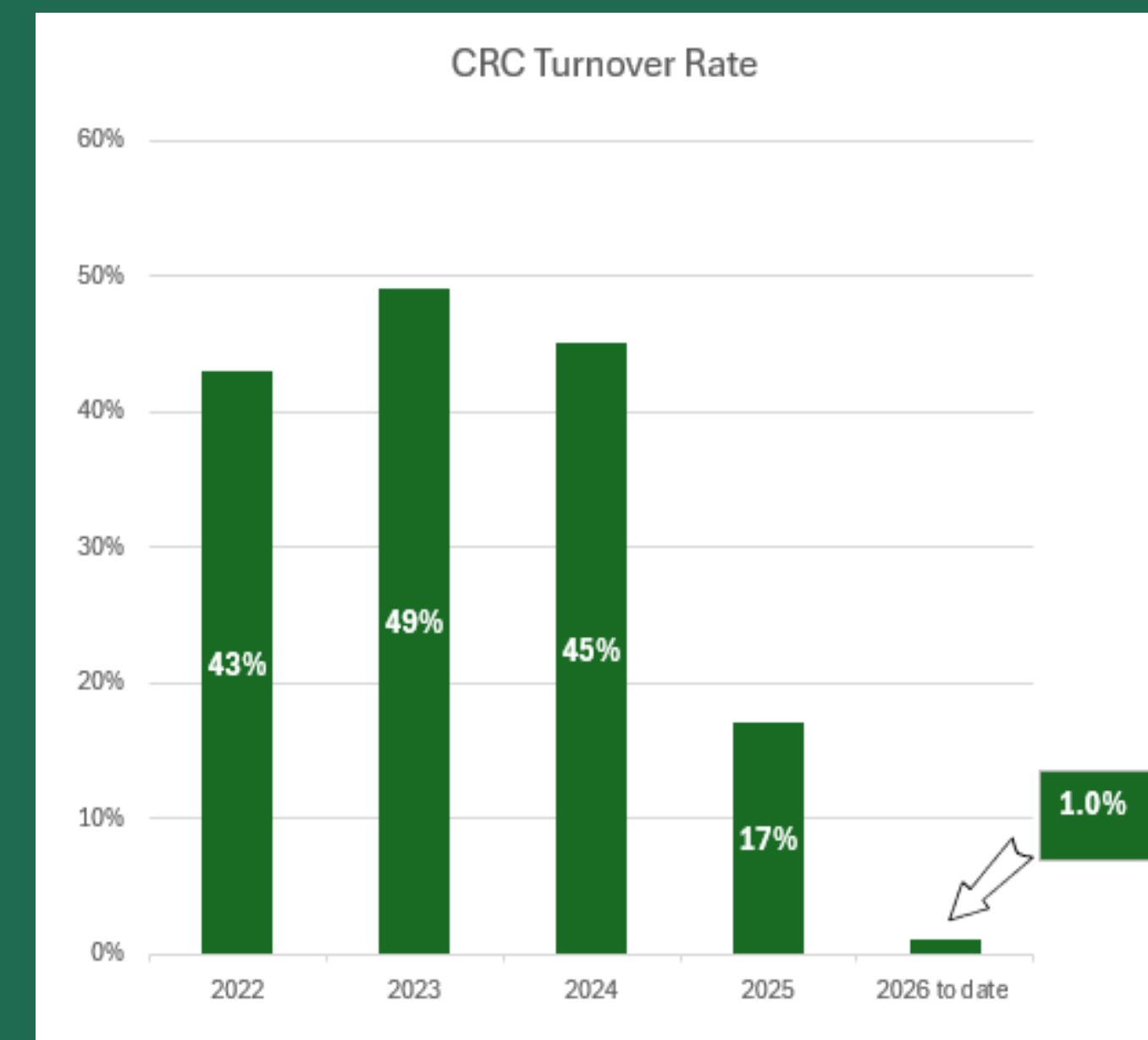
Solutions

In alignment with broader optimization initiatives including workload assessment, workflow redesign, and study pipeline management, our Center implemented structured onboarding, competency-based training, annual competency evaluation, and ongoing professional development.

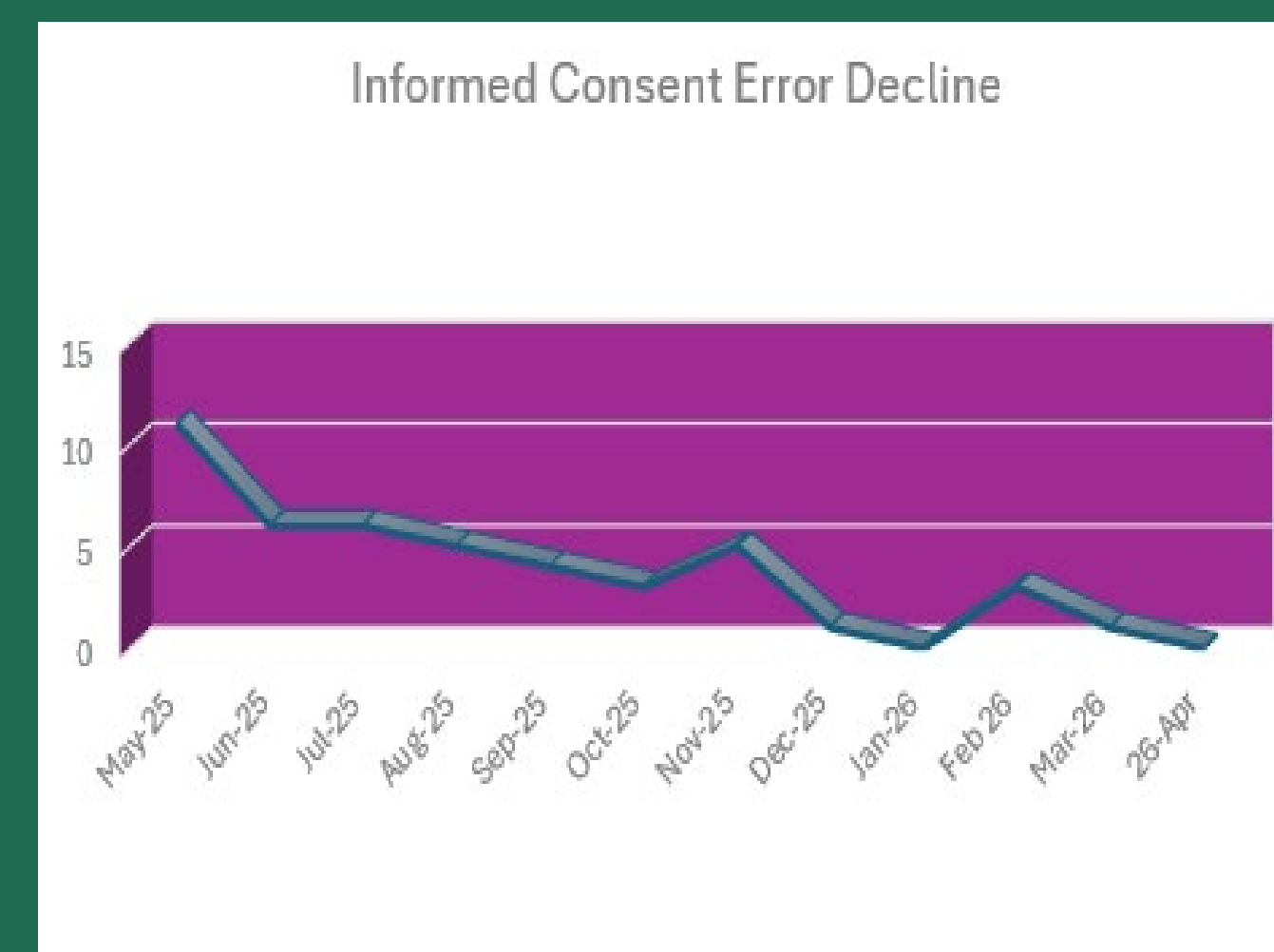
Disclosure Information

None

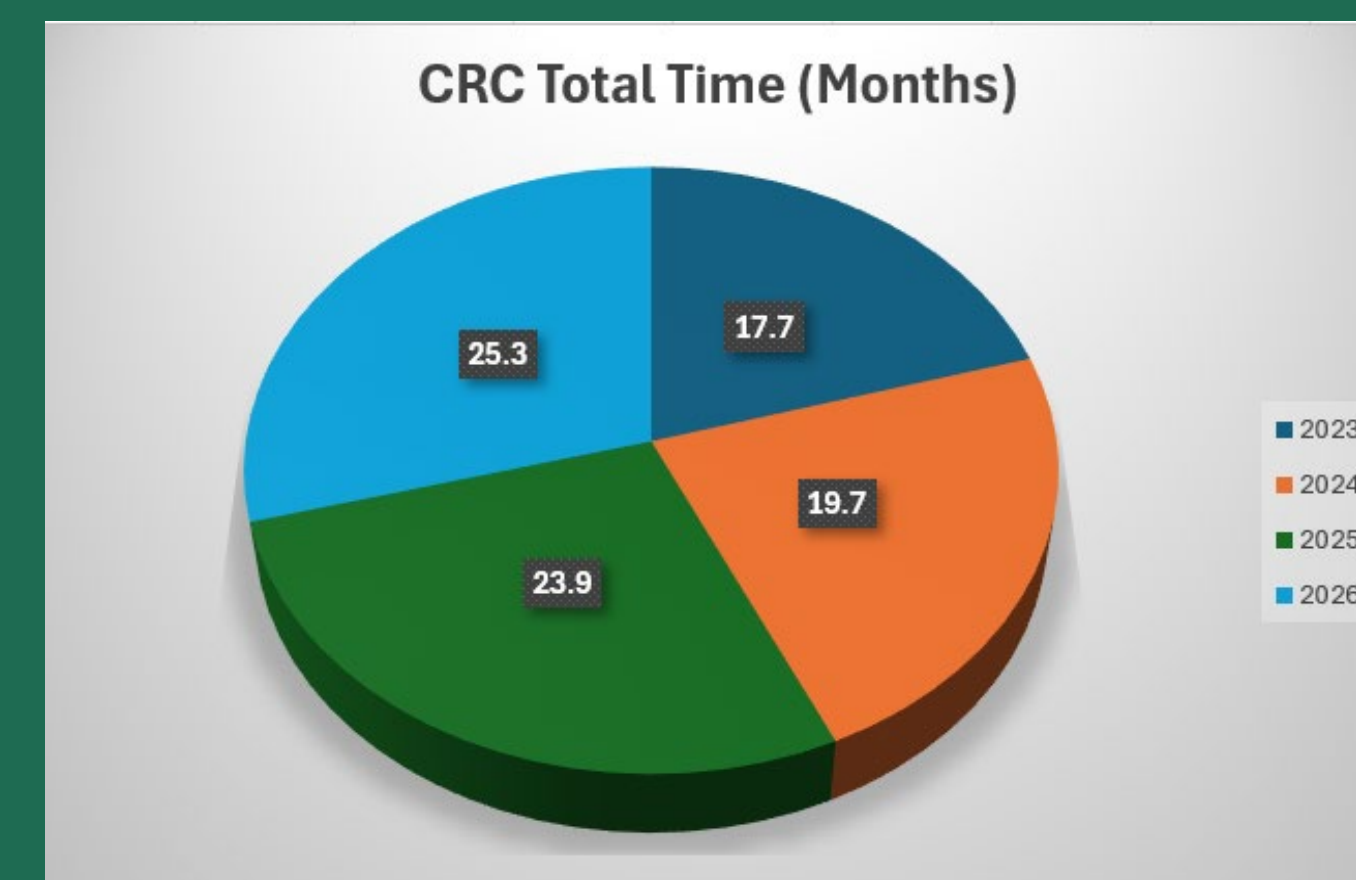
Outcomes



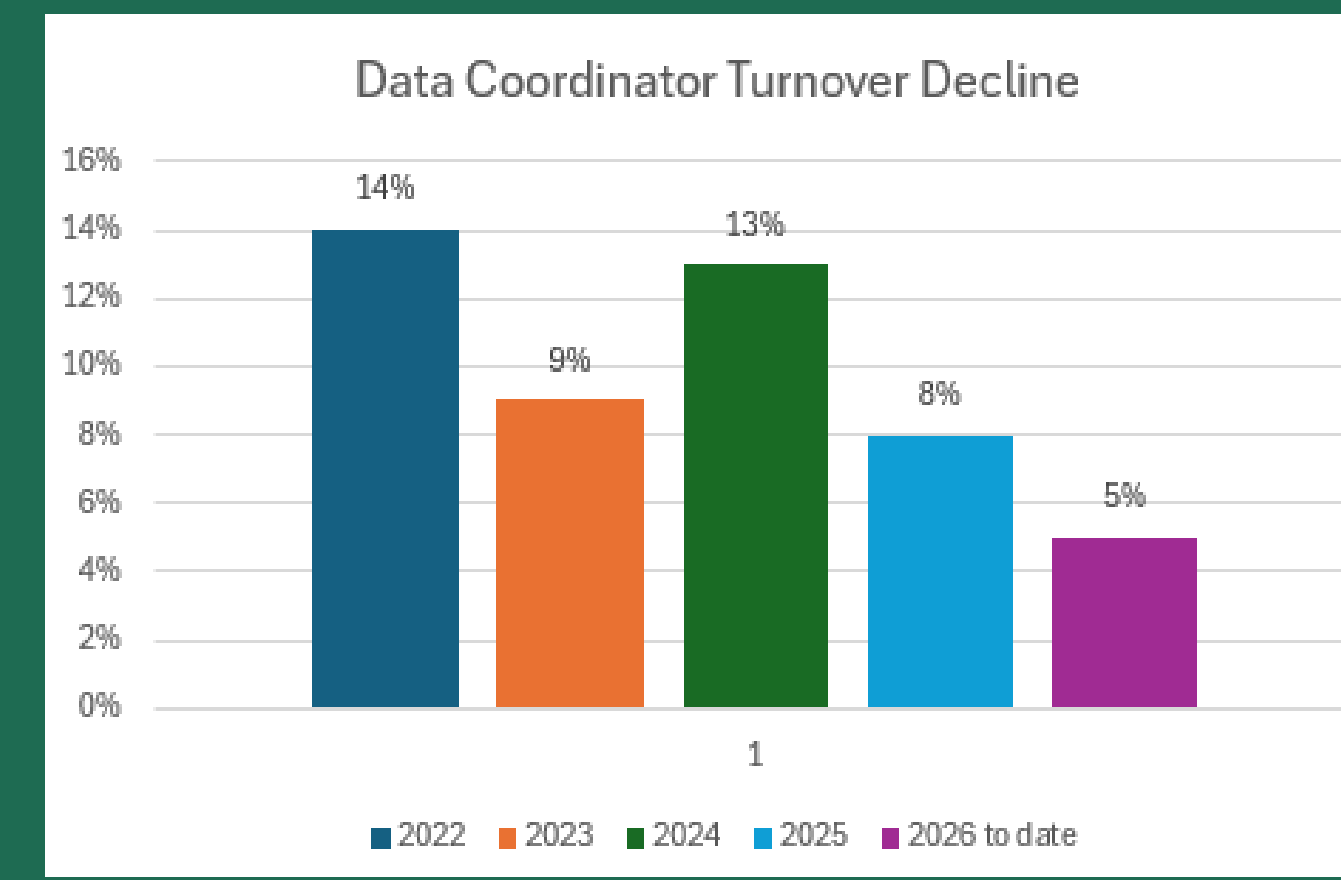
The turnover rate of CRC has dramatically declined- from 49 percent in 2023 to 17 percent in 2025 as our training program has fully developed within the CRC role.



2025 tracking of informed consent errors showed a steady decline as real time review was implemented



For CRCs that have resigned from the CTO since structured training has been implemented, the length of employment continues to increase each year



Turnover rate of DC has declined- from 14 percent in 2022 to 8 percent in 2025. 2026 will hopefully remain at a steady decline.

Future Directions

Future directions include expanding the recently implemented research training for investigators, nursing, and ancillary staff through the Cancer Service Line and further expanding our educational footprint in the inpatient setting to continue our efforts to seamlessly integrate clinical research operations into standard of care processes at UAB.

Methods

Twelve-week initial onboarding includes in person lessons, self-guided electronic training modules and assessments, and role specific shadowing with research duties performed under supervision. Dedicated paper-based competencies assess readiness for independent work.

A sixth month post onboarding assessment is completed to evaluate learning outcomes presented in weeks one to twelve.

All staff complete an annual assessment and competencies evaluating quality of work. Results are utilized for staff feedback on performance evaluation as well as readiness for promotion.

Continued education is provided at least once monthly delivered in person and distance accessible focusing on broader research topics.

These initiatives strengthen staff proficiency in Good Clinical Practice (GCP), protocol execution, and regulatory compliance and foster a supportive work environment that promotes engagement and professional growth.

Lessons Learned

Structured training assists in stabilizing staff turnover. Training content must be tailored to the career ladder level, experience, and medical and clinical research knowledge of the staff. Providing data-based feedback to the staff on their strengths and weaknesses builds trust and allows assignment of acuity of trials to match the knowledge level of the employee. An on-site education team provides an additional layer of real-time support for management and staff, promoting a culture of collaboration. While we recognize the requirements for training may seem overwhelming, our data supports the effectiveness of the program.